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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/616,101	07/08/2003	Ying Luo	RIGL-010CIP3	5361
24353	7590 01/19/2006	EXAMINER		
BOZICEVIC, FIELD & FRANCIS LLP 1900 UNIVERSITY AVENUE			RAO, MANJUNATH N	
SUITE 200				PAPER NUMBER
EAST PALO	O ALTO, CA 94303		1652	

DATE MAILED: 01/19/2006

Please find below and/or attached an Office communication concerning this application or proceeding.

,	Application No.	Applicant(s)
	10/616,101	LUO ET AL.
Office Action Summary	Examiner	Art Unit
	Manjunath N. Rao, Ph.D.	1652
The MAILING DATE of this communication ap Period for Reply	pears on the cover sheet with the c	correspondence address
A SHORTENED STATUTORY PERIOD FOR REPL WHICHEVER IS LONGER, FROM THE MAILING D.  - Extensions of time may be available under the provisions of 37 CFR 1. after SIX (6) MONTHS from the mailing date of this communication.  - If NO period for reply is specified above, the maximum statutory period.  - Failure to reply within the set or extended period for reply will, by statut Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	DATE OF THIS COMMUNICATION 136(a). In no event, however, may a reply be tir will apply and will expire SIX (6) MONTHS from e, cause the application to become ABANDONE	N. nely filed the mailing date of this communication. D (35 U.S.C. § 133).
Status		
1)⊠ Responsive to communication(s) filed on 31 ⊆ 2a)□ This action is <b>FINAL</b> . 2b)□ Thi 3)□ Since this application is in condition for allowed closed in accordance with the practice under	s action is non-final. ance except for formal matters, pro	
Disposition of Claims		
4) ☐ Claim(s) <u>1-37</u> is/are pending in the application 4a) Of the above claim(s) is/are withdra 5) ☐ Claim(s) is/are allowed. 6) ☐ Claim(s) is/are rejected. 7) ☐ Claim(s) is/are objected to. 8) ☐ Claim(s) <u>1-37</u> are subject to restriction and/or	awn from consideration.	
Application Papers		
9) The specification is objected to by the Examina 10) The drawing(s) filed on is/are: a) accomposed and applicant may not request that any objection to the Replacement drawing sheet(s) including the correct and the specific production is objected to by the Examination is objected to by the Examination is objected.	cepted or b) objected to by the drawing(s) be held in abeyance. Section is required if the drawing(s) is ob	e 37 CFR 1.85(a). jected to. See 37 CFR 1.121(d).
Priority under 35 U.S.C. § 119		
12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of:  1. Certified copies of the priority documen 2. Certified copies of the priority documen 3. Copies of the certified copies of the priority application from the International Burea * See the attached detailed Office action for a list	nts have been received. Its have been received in Applicationity documents have been received in Application (PCT Rule 17.2(a)).	ion No ed in this National Stage
Attachment(s)	<b></b>	
<ol> <li>Notice of References Cited (PTO-892)</li> <li>Notice of Draftsperson's Patent Drawing Review (PTO-948)</li> <li>Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08 Paper No(s)/Mail Date</li> </ol>	4)  Interview Summary Paper No(s)/Mail D  5)  Notice of Informal F  6) Other:	

## **DETAILED ACTION**

Claims 1-37 are currently pending this application.

## Election/Restrictions

Restriction to one of the following inventions is required under 35 U.S.C. 121:

- I. Claim 1, drawn to a method for screening for a bioactive agent capable of binding to a cell cycle protein tankyrase H, classified in class 435, subclass 4.
- II. Claims 2-3, drawn to a method for screening for agents capable of interfering with the binding of tankyrase H with p21 protein, classified in class 435, subclass 4.
- III. Claims 4-5, 7-8 drawn to a method for screening for a bioactive agent capable of modulating the activity of tankyrase H, classified in class 435, subclass 4.
  (Please note that claim 8 improperly depends on claim 25 which is actually drawn to an antibody. However, Examiner has included claim 8 in the above group as it appears that claim should depend from claim 7).
- IV. Claim 6, drawn to a method of diagnosing cancer by determining the activity of tankyrase, classified in class 435, subclass 15.
- V. Claim 9, drawn to a method of treating an individual with a cell cycle related disorder using the inhibitor of TaHo, classified in class 514, subclass 789.
- VI. Claims 10-19, drawn to polynucleotides encoding a protein, a expression vector, a host cell and a process of producing the protein, classified in class 435, subclass 69.1.

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VII. Claims 20-23, drawn to a recombinant polypeptide with SEQ ID NO:3 or 4, classified in class 435, subclass 196.

- VIII. Claims 24-26, drawn to an antibody, classified in class 530, subclass 387.5.
- IX. Claim 27-30, drawn to a method of screening for a bioactive agent capable of modulating PARP activity, classified in class 435, subclass 19.
- X. Claims 31-34, drawn to a method of screening a candidate agent capable of inhibiting proliferation, classified in class 435, subclass 325.
- XI. Claims 35-37, drawn to a method for inhibiting growth of a tumor cell, classified in class 435, subclass 344.

The inventions are distinct, each from the other because of the following reasons:

Inventions I through V, IX through XI are patentably distinct from each other. The method of group I drawn to screening for bioactive compounds capable of binding to tankyrase H, the method of group II drawn to screening for agents which interfere in binding to protein p21, the method of group III drawn to screening for agents which modulate the activity of tankyrase H, the method of group IV drawn to diagnosing cancer, the method of group V drawn to treating an individual with a cell cycle disorder, the method of group IX drawn to screening for bioactive agent capable of modulating PARP activity, the method of group X drawn to screening a candidate agent capable of inhibiting proliferation and the method of group XI drawn to inhibiting growth of a tumor cell are all unrelated as they comprise distinct steps, utilize different products and produce different results. The groups have acquired separate status in the art and separate fields of search as further evidenced by their separate classification.

Inventions VI, VII, VIII are patentably distinct from each other. The polynucleotide of group VI, the polypeptide of group VII, and the antibody of group VIII, each comprise amino acid sequences and nucleotide sequences which are chemically unrelated, do not require each other for practice; have separate utilities, such as use of the group VII polypeptide to regulate cell cycle versus the use of polynucleotide in a hybridization reaction, versus the use of the antibody in an affinity purification reaction and are subject to separate manufacture and sale. The groups have acquired separate status in the art and separate fields of search.

Inventions VII and I through IV, IX, X are related as product and process of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product (MPEP § 806.05(h)). In the instant case the polypeptide of Group VII can be used to raise specific antibodies as opposed to its use in the methods of Groups I through IV, IX, X.

Inventions VIII and V, XI are related as product and process of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product (MPEP § 806.05(h)). In the instant case the antibody of Group VIII can be used to affinity purify the polypeptide as opposed to its use in the methods of Groups V, XI.

Because these inventions are distinct for the reasons given above and have acquired a separate status in the art as shown by their different classification, restriction for examination purposes as indicated is proper.

Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

## Rejoinder of restricted inventions

The examiner has required restriction between product and process claims. Where applicant elects claims directed to the product, and a product claim is subsequently found allowable, withdrawn process claims that depend from or otherwise include all the limitations of the allowable product claim will be rejoined in accordance with the provisions of MPEP § 821.04. Process claims that depend from or otherwise include all the limitations of the patentable product will be entered if the amendment is presented prior to final rejection or allowance, whichever is earlier. Amendments submitted after final rejection are governed by 37 CFR 1.116; amendments submitted after allowance are governed by 37 CFR 1.312. In the event of rejoinder, the requirement for restriction between the product claims and the rejoined

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process claims will be withdrawn, and the rejoined process claims will be fully examined for patentability in accordance with 37 CFR 1.104. Thus, to be allowable, the rejoined claims must meet all criteria for patentability including the requirements of 35 U.S.C. 101, 102, 103, and 112. Until an elected product claim is found allowable, an otherwise proper restriction requirement between product claims and process claims may be maintained. Withdrawn process claims that are not commensurate in scope with an allowed product claim will not be rejoined. See "Guidance on Treatment of Product and Process Claims in light of *In re Ochiai, In re Brouwer* and 35 U.S.C. § 103(b)," 1184 O.G. 86 (March 26, 1996). Additionally, in order to rejoin, in accordance with the above policy, applicant is advised that the process claims should be amended during prosecution either to maintain dependency on the product claims or to otherwise include the limitations of the product claims. Failure to do so may result in a loss of rejoinder. Further, note that the prohibition against double patenting rejections of 35 U.S.C. 121 does not apply where the restriction requirement is withdrawn by the examiner before the patent issues. See MPEP § 804.01.

Any inquiry concerning this communication or earlier communications from the Examiner should be directed to Manjunath N. Rao, Ph.D. whose telephone number is 571-272-0939. The Examiner can normally be reached on 7.00 a.m. to 3.30 p.m. If attempts to reach the examiner by telephone are unsuccessful, the Examiner's supervisor, Ponnathapura Achutamurthy can be reached on 571-272-0928. The fax phone numbers for the organization where this application or proceeding is assigned is 571-273-8300 for regular communications and for After Final communications.

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Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is 571-272-1600.

Manjunath N. Rao, Ph.D.

Primary Examiner Art Unit 1652

January 7, 2006